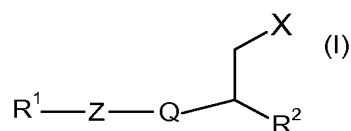


### Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

What is claimed is:

1. **(Currently Amended)** A compound of formula (I):



wherein:

R<sup>1</sup> is optionally substituted -C<sub>4-12</sub> alkyl, -C<sub>2-10</sub>alkylcycloalkyl, C<sub>2-6</sub>alkylheterocycloalkyl, -C<sub>2-6</sub>alkylaryl, optionally substituted 5- or 6- membered aryl or heteroaryl with the proviso that R<sup>2</sup> is not pyridinyl;

Z is a bond, CH<sub>2</sub>, O, S, SO, SO<sub>2</sub>, NR<sup>4</sup>, OCR<sup>4</sup>R<sup>5</sup> or CR<sup>4</sup>R<sup>5</sup>O; or Z, R<sup>1</sup> and Q together form an optionally substituted fused tricyclic group;

Q is an optionally substituted 5- or 6- membered aryl or heteroaryl ring;

X is COR<sup>3</sup>;

R<sup>2</sup> is CONH<sub>2</sub>, CO<sub>2</sub>H, CO<sub>2</sub>R<sup>7</sup>, SO<sub>2</sub>R<sup>7</sup> or SO<sub>2</sub>NR<sup>8</sup>R<sup>9</sup>, \_\_\_\_\_  
**with the proviso provided** that R<sup>2</sup> is not CO<sub>2</sub>R<sup>7</sup>, when X is CONH<sub>2</sub>;

R<sup>3</sup> is OR<sup>6</sup> or NR<sup>8</sup>R<sup>9</sup>;

R<sup>4</sup> and R<sup>5</sup> each independently is H, C<sub>1-6</sub> alkyl or C<sub>1-4</sub> alkylaryl;

R<sup>6</sup> is H or C<sub>1-6</sub> alkyl;

R<sup>7</sup> is C<sub>1-6</sub> alkyl; and

R<sup>8</sup> and R<sup>9</sup> each independently is H or C<sub>1-6</sub> alkyl; or R<sup>8</sup> and R<sup>9</sup> together with the nitrogen atom to which they are attached form a 5- or 6- membered ring which may optionally include 1 or more further heteroatoms selected from O, S and N;  
or

physiologically functional derivatives thereof,

**with the proviso provided** that formula (I) compounds are not:

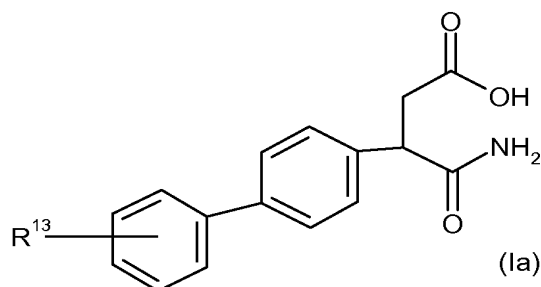
[3-(acetylamino)-4-cyclohexylphenyl]-butanedioic acid and 3-(acetylamino)-4-cyclohexylphenyl]-butanedioic acid diethyl ether;

butanedioic acid [3-methoxy-4-(phenylmethoxy)phenyl]; or

butanedioic acid [4-(phenylmethoxy)phenyl]; and

**with the proviso further provided** that when R<sup>1</sup> is C<sub>4-12</sub>alkyl, Z is other than a bond, O or CH<sub>2</sub>, or physiologically functional derivatives thereof.

2. (Previously Presented) A compound as claimed in claim 1 wherein X is CO<sub>2</sub>H and R<sup>2</sup> represents CONH<sub>2</sub>.
3. (Previously Presented) A compound as claimed in claim 1 wherein Q is an unsubstituted phenyl.
4. (Previously Presented) A compound as claimed in claim 1 wherein Z represents a bond or O.
5. (Previously Presented) A compound as claimed in claim 1 of formula (Ia):

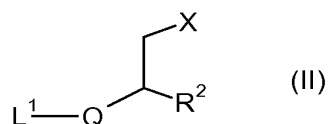


wherein R<sup>13</sup> is H, halo, CF<sub>3</sub>, -OCF<sub>3</sub>, cyano, nitro, OR<sup>14</sup>, SR<sup>15</sup> or COR<sup>16</sup>; and R<sup>14</sup>, R<sup>15</sup>, R<sup>16</sup> independently are H, C<sub>1-6</sub> alkyl or C<sub>1-4</sub> alkylaryl; or physiologically functional derivatives thereof.

6. (Cancelled)
7. (Previously Presented) A method for the treatment of a human or animal subject suffering from or susceptible to an inflammatory disease or an autoimmune disorder which method comprises administering to said subject an effective amount of a compound as claimed in claim 1.
8. (Cancelled)
9. (Previously Presented) A pharmaceutical composition comprising a compound as claimed in claim 1 and a pharmaceutically acceptable carrier.

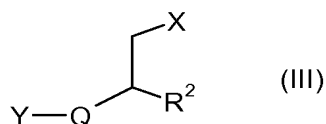
10. **(Currently Amended)** A process for ~~the~~ preparation of compounds of formula (I) as defined in claim 1, ~~which~~ wherein the process comprises:

(A) ~~for~~ preparing a compound of formula (I), wherein Z is a bond and R<sup>1</sup> is an optionally substituted 5- or 6- membered aryl or heteroaryl, by reacting a compound of formula (II):



wherein R<sup>2</sup>, Q and X are as previously defined for formula (I) and L<sup>1</sup> is a leaving group, with a reagent suitable to introduce the group R<sup>1</sup>; or

(B) (i) ~~for~~ preparing a compound of formula (I), wherein Z is O, S, SO, SO<sub>2</sub>, NR<sup>4</sup> or OCR<sup>4</sup>R<sup>5</sup>, by reacting a compound of formula (III):



wherein R<sup>2</sup>, Q and X are as previously defined for formula (I) and Y is OH, SH, NHR<sup>4</sup> or HOOCR<sup>4</sup>R<sup>5</sup>, with a compound of formula (IV):



wherein R<sup>1</sup> is defined above for compounds of formula (I) and L<sup>2</sup> represents a leaving group; and

(ii) wherein Y is  $\text{---SH}$ , optionally followed by ~~oxidation~~ oxidizing the Y group to the corresponding SO or SO<sub>2</sub> group as required; or

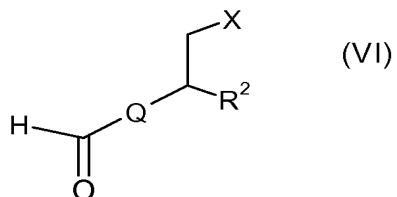
(C) ~~for~~ preparing a compound of formula (I), wherein Z is  $\text{---CR}^4\text{R}^5\text{O---}$ , by reacting a compound of formula (III), wherein Y is  $\text{---OH}$ , with a compound of formula (V):



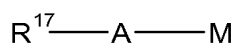
wherein  $R^1$ ,  $R^4$ ,  $R^5$  are defined above for compounds of formula (I) and  $L^3$  represents a leaving group; or

(D) ~~for~~ preparing a compound of formula (I), wherein Z is  $CH_2$  and  $R^1$  is an optionally substituted 5- or 6- membered aryl or heteroaryl, by reacting

\_\_\_\_\_ (i) a compound of formula (VI):



wherein Q, X and  $R^2$  are as defined above, with an optionally substituted 5- or 6- membered aryl or heteroaryl nucleophile, which is a compound of formula (VII):



(VII)

wherein A is a 5- or 6- membered aryl or heteroaryl,  $R^{17}$  is H or one or more substituents and M is a metal and

\_\_\_\_\_ (ii) ~~reduction and elimination~~ reducing and eliminating of the a resultant or product alcohol formed from step (i); and, if necessary,

(E) ~~deprotection~~ optionally deprotecting of a protected form of compounds of formula (I) with a protecting group.